

palladium, *Borrelia burgdorferi*, *Borrelia recurrentis*, *Rickettsial* pathogens, *Nocardia*, and *Acitnomycetes*.

Fungal infectious agents which can be detected by the present invention include *Cryptococcus neoformans*, *Blastomyces dermatitidis*, *Histoplasma*
5 *capsulatum*, *Coccidioides immitis*, *Paracoccidioides brasiliensis*, *Candida albicans*, *Aspergillus fumigautus*, *Phycomycetes* (*Rhizopus*), *Sporothrix schenckii*, *Chromomycosis*, and *Maduromycosis*.

Viral infectious agents which can be detected by the present invention include human immunodeficiency virus, human T-cell lymphocytotropic virus,
10 hepatitis viruses (e.g., Hepatitis B Virus and Hepatitis C Virus), Epstein-Barr Virus, cytomegalovirus, human papillomaviruses, orthomyxo viruses, paramyxo viruses, adenoviruses, corona viruses, rhabdo viruses, polio viruses, toga viruses, bunya viruses, arena viruses, rubella viruses, and reo viruses.

Parasitic agents which can be detected by the present invention include
15 *Plasmodium falciparum*, *Plasmodium malaria*, *Plasmodium vivax*, *Plasmodium ovale*, *Onchoverva volvulus*, *Leishmania*, *Trypanosoma* spp., *Schistosoma* spp., *Entamoeba histolytica*, *Cryptosporidium*, *Giardia* spp., *Trichimonas* spp., *Balatidium coli*, *Wuchereria bancrofti*, *Toxoplasma* spp., *Enterobius vermicularis*, *Ascaris lumbricoides*, *Trichuris trichiura*, *Dracunculus medinesis*, trematodes,
20 *Diphyllobothrium latum*, *Taenia* spp., *Pneumocystis carinii*, and *Necator americanis*.

The present invention is also useful for detection of drug resistance by infectious agents. For example, vancomycin-resistant *Enterococcus faecium*, methicillin-resistant *Staphylococcus aureus*, penicillin-resistant *Streptococcus pneumoniae*, multi-drug resistant *Mycobacterium tuberculosis*, and AZT-resistant
25 human immunodeficiency virus can all be identified with the present invention.

Genetic diseases can also be detected by the process of the present invention. This can be carried out by prenatal or post-natal screening for chromosomal and genetic aberrations or for genetic diseases. Examples of detectable genetic diseases include: 21 hydroxylase deficiency, cystic fibrosis, Fragile X
30 Syndrome, Turner Syndrome, Duchenne Muscular Dystrophy, Down Syndrome or other trisomies, heart disease, single gene diseases, HLA typing, phenylketonuria, sickle cell anemia, Tay-Sachs Disease, thalassemia, Klinefelter Syndrome,

Huntington Disease, autoimmune diseases, lipodosis, obesity defects, hemophilia, inborn errors of metabolism, and diabetes.

Cancers which can be detected by the process of the present invention generally involve oncogenes, tumor suppressor genes, or genes involved in DNA amplification, replication, recombination, or repair. Examples of these include: BRCA1 gene, p53 gene, APC gene, Her2/Neu amplification, Bcr/Ab1, K-ras gene, and human papillomavirus Types 16 and 18. Various aspects of the present invention can be used to identify amplifications, large deletions as well as point mutations and small deletions/insertions of the above genes in the following common human cancers: leukemia, colon cancer, breast cancer, lung cancer, prostate cancer, brain tumors, central nervous system tumors, bladder tumors, melanomas, liver cancer, osteosarcoma and other bone cancers, testicular and ovarian carcinomas, head and neck tumors, and cervical neoplasms.

In the area of environmental monitoring, the present invention can be used for detection, identification, and monitoring of pathogenic and indigenous microorganisms in natural and engineered ecosystems and microcosms such as in municipal waste water purification systems and water reservoirs or in polluted areas undergoing bioremediation. It is also possible to detect plasmids containing genes that can metabolize xenobiotics, to monitor specific target microorganisms in population dynamic studies, or either to detect, identify, or monitor genetically modified microorganisms in the environment and in industrial plants.

The present invention can also be used in a variety of forensic areas, including for human identification for military personnel and criminal investigation, paternity testing and family relation analysis, HLA compatibility typing, and screening blood, sperm, or transplantation organs for contamination.

In the food and feed industry, the present invention has a wide variety of applications. For example, it can be used for identification and characterization of production organisms such as yeast for production of beer, wine, cheese, yogurt, bread, etc. Another area of use is with regard to quality control and certification of products and processes (e.g., livestock, pasteurization, and meat processing) for contaminants. Other uses include the characterization of plants, bulbs, and seeds for

breeding purposes, identification of the presence of plant-specific pathogens, and detection and identification of veterinary infections.

Desirably, the oligonucleotide probes are suitable for ligation together at a ligation junction when hybridized adjacent to one another on a corresponding
5 target nucleotide sequence due to perfect complementarity at the ligation junction. However, when the oligonucleotide probes in the set are hybridized to any other nucleotide sequence present in the sample, there is a mismatch at a base at the ligation junction which interferes with ligation. Most preferably, the mismatch is at the base at the 3' base at the ligation junction. Alternatively, the mismatch can be at the bases
10 adjacent to bases at the ligation junction.

Before carrying out the ligase detection reaction, in accordance with the present invention, target nucleotide sequences in the sample can be preliminarily amplified. This preferably carried out with polymerase chain reaction. The polymerase chain reaction process is fully described in H. Erlich, et. al., "Recent
15 Advances in the Polymerase Chain Reaction," Science 252: 1643-50 (1991); M. Innis, et. al., PCR Protocols: A Guide to Methods and Applications, Academic Press: New York (1990) and R. Saiki, et. al., "Primer-directed Enzymatic Amplification of DNA with a Thermostable DNA Polymerase," Science 239: 487-91 (1988), which are hereby incorporated by reference.

20 The ligase detection reaction process achieves linear amplification of a target sequence. The ligase chain reaction utilizes essentially the same steps and conditions as the ligase detection reaction to achieve exponential amplification. This greater level of amplification is achieved by utilizing 2 sets of complementary oligonucleotides with each set hybridizing to complementary strands of a target
25 nucleic acid sequence.

In the ligase chain reaction process of the present invention, the presence of a target double stranded nucleic acid formed from first and second complementary target nucleotide sequences is detected in a sample. The target double stranded nucleic acid differs from other nucleotide sequences by one or more single
30 base changes, insertions, deletions, or translocations.

This method involves providing a sample potentially containing a target double stranded nucleic acid formed from first and second complementary